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EDGE ARTICLE

Surface plasmon resonance imaging of glycoarrays identifies novel and unnatural carbohydrate-based ligands for potential ricin sensor development†

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Carbohydrate microarrays provide access to high through-put analysis of protein–carbohydrate interactions. Herein we demonstrate the use of SPR imaging (SPRi) of glycoarrays to assess the ligand specificity of the reputedly galactose-specific plant lectin RCA₁₂₀ (*Ricinus communis* agglutinin 120), a surrogate for the bioterrorism agent ricin. Glycoarray studies identified RCA₁₂₀ ligands based on galactose substituted at the 6-position with sialic acid. These observations, which were confirmed by saturation transfer difference (STD) NMR spectroscopy studies, inspired the synthesis of non-natural 6-substituted galactose derivatives, which were shown to have ~3–4 fold enhanced binding to RCA₁₂₀ with respect to the unsubstituted compound. These novel unnatural galactosides, which are chemically and biologically more robust than their natural glycan counterparts, represent new potential ligands for the development of carbohydrate-based ricin sensors.

Introduction

Major advances in analytical glycoscience over the past decade or so present many new opportunities for identifying carbohydrate–ligand interactions of biological significance.^{1–4} In particular, the development of carbohydrate and lectin microarrays^{5–9} provides access to quantitative, high throughput screening approaches that could not previously have been contemplated. In addition to the identification of natural ligands, these technologies also provide a basis for the rapid optimisation of non-natural ligands for carbohydrate-binding proteins, as exemplified by work on novel sialic acid derivatives,¹⁰ some of which serve as high affinity ligands for sialic acid-binding lectins (Siglecs).¹¹ In the course of work on the development of ricin sensors,¹² we had cause to consider the identification of novel ligands, and hence turned to carbohydrate microarrays.¹³

Ricin is a lethal, ribosome-inactivating protein^{14,15} found in plants that are used for castor oil production. Its lectin-like B-chain facilitates adhesion of the toxin to non-reducing terminal galactose residues of glycoproteins on mammalian cell surfaces,

resulting in the delivery of the toxic A-chain warhead into the cytosol. The A-chain catalyses the hydrolytic cleavage of a single base from eukaryotic ribosomal RNA, leading to a shutdown in protein synthesis and ultimately cell death. Ricin is several thousand times more toxic to man than cyanide, with the lethal dose for an adult estimated to be around 500 ng Kg^{−1} of body weight.^{14,15} Due to its high toxicity and accessibility, its potential as a biowarfare agent has long been recognised, with several recent incidents in the UK and the US resulting in threats to public safety.^{14,15} While recent efforts have identified potential leads for the development of anti-ricin inhibitors,¹⁶ a rapid method for the early detection of traces of the toxin remains an important goal.¶ Given the carbohydrate-binding properties of ricin B-chain, carbohydrate-based sensors are potentially attractive. Certain systems have been shown to exhibit very high selectivity for bacterial toxins¹⁷ and plant lectins, including ricin;^{12,18,19} related carbohydrate-based sensing platforms have also been used for bacterial detection.^{20–25} In addition, galactose-coated gold nanoparticles in a colorimetric assay¹² for the detection of *Ricinus communis* agglutinin (RCA₁₂₀), a surrogate for ricin (RCA₆₀),²⁶ afforded higher sensitivity than antibody-conjugated nanoparticles in piezoelectric immunosensing.²⁷

Despite these early developments, D-galactosides are prevalent in nature and many other mammalian, plant and microbial lectins and enzymes recognise and/or metabolise this common monosaccharide.²⁸ Selectivity is therefore a primary issue. In the context of robust ricin sensor development, we were drawn to explore carbohydrate microarrays in the search for alternative naturally occurring ligands for ricin that might provide insight for the design of non-natural ligands that in turn could show

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improved selectivity. We noted that surface plasmon resonance approaches had already been used successfully in conjunction with D-galactoside surfaces to develop biosensors for ricin.^{18,19} With a view to exploring a much wider range of potential toxin ligands, we were drawn to explore surface plasmon resonance imaging (SPRi), an array-based approach that is finding wide use in bimolecular interaction analysis and for the development of affinity-based biosensors.²⁹ We have recently employed SPRi for the analysis of protein interactions with carbohydrate microarrays.¹³ Here we report the further use of this technique to analyse the interaction of RCA₁₂₀ with an array of natural glycans, which in turn informed studies on ligand (re)design to provide unnatural sugar-based ligands for potential use in ricin sensors.

Results and discussion

Analysis of RCA₁₂₀ interaction with a carbohydrate microarray by SPR imaging

Array generation through split-pin, contact printing of a library of 40 biotinylated glycans (Table 1) onto commercial neutravidin-coated gold sensor chips was carried out as reported previously.¹³ To briefly summarise, the library assessed was composed of a range of glycans with non-reducing termini based on: Gal-β-1,4 (7 compounds), Gal-β-1,3 (3), GalNAc (1), GlcNAc (4), sulfo-Gal (3), NeuAc-2,3 (12), NeuGc-2,3/2,6 (2/1), NeuAc-2,6 (4), and NeuAc-2,8 (3) (Table 1). The array was used to screen for RCA₁₂₀ binders in the anticipation that glycans possessing a non-reducing terminal β-linked galactose residue (black and orange), the previously determined simple ligand for this lectin, would stand out as hits.^{13,18,19} Data for equilibrium binding of RCA₁₂₀ and the α-2,6-NeuAc-specific *Sambucus nigra* agglutinin (SNA) to the glycoarray are presented in Fig. 1.*

It was immediately evident from these array data that RCA₁₂₀ had a preference for binding to non-reducing β-D-galactose (strongest responses are black and orange bars). However, such recognition was context-dependent, with Galβ1-4GlcNAc (B81) giving a stronger response than Galβ1-4Glc (B80), for instance; an extended glycan chain (e.g. B115) also gave a stronger response than the shorter counterpart (B81). Some galactose-terminated structures were relatively poorly recognised, in particular branched glycans, such as Le^x trisaccharide (B266) and Le^a trisaccharide (B156). As expected, RCA₁₂₀ did not bind significantly to sugars with non-reducing terminal GlcNAc (pink) but results with sialic acid (NeuAc)-containing ligands were intriguing (red, green, blue, purple bars). On the whole, glycans terminated in α-2,3Neu5Ac/Gc (red bars, plus B89, B90) were poor ligands whilst those terminated in α-2,6Neu5Ac/Gc (blue bars, plus B93) showed relatively good binding to RCA₁₂₀. Extending the α-2,3Neu5Ac arm with an additional α-2,8Neu5Ac residue did not recover binding (purple bars). These data are consistent with the ability of RCA₁₂₀ to accommodate substitution at the 6-position of the non-reducing terminal β-galactose with sialic acid (Fig. 2); likewise, terminal galactose 6-sulfate was recognised by RCA₁₂₀ whereas galactose 3-sulfate was not (cyan bars, B185 and B197, respectively). Together these data comprehensively map novel functional ‘hotspots’ for RCA₁₂₀ carbohydrate binding.

Analysis of SNA interaction with a carbohydrate microarray by SPR imaging

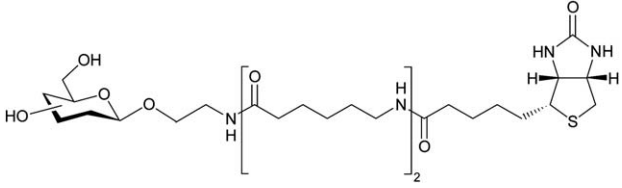
Given these unexpected interactions of RCA₁₂₀ with α-2,6-NeuAc-containing glycans, the glycoarray was probed with another previously characterised α-2,6-NeuAc-specific lectin, in this case *Sambucus nigra* agglutinin (SNA) (Fig. 1B). For SNA, the only glycan ligands that were identified in this study were three of the four α-2,6-NeuAc compounds (B87, B179, B274) and the single α-2,6-NeuGc (B93) out of the library of forty glycans. Together, these data confirmed the functional performance of the array bearing this key ligand type.

Development of novel carbohydrate-based ligands for RCA₁₂₀

The novel glycoarray data above revealed 6-modified D-galactosides as good RCA₁₂₀ ligands. To explore the potential of these observations in the creation of bespoke ricin ligands a series of Galβ1,4-GlcNAc derivatives modified at the galactose 6-position were designed. The intention here was to attempt to move away from the potentially biologically and chemically labile α-2,6-linked NeuAc compounds in favour of a more robust ligand, which would be necessary for sensor applications. A library of Galβ1,4-GlcNAc derivatives modified at the galactose 6-position (1–7) was devised (Scheme 1), with a view to post-synthesis biotinylation through hydrazone chemistry^{32,33} (giving 1Bt–7Bt) to enable SPRi array analysis, as above.

Syntheses of target derivatives employed the coupling of appropriately tailored glycosyl trichloroacetimidate donors (8–12) with a common partially protected GlcNAc acceptor (13). Known trichloroacetimidates³⁴ were prepared from five variously modified galactose derivatives: D-galactose (8, 97% yield), 6-deoxy-6-fluoro-D-galactose (9),³⁵ D-fucose (10)³⁶ (6-deoxy-D-galactose), arabinose (11)³⁷ (deshydroxymethyl-D-galactose) and 6-deoxy-6-azido-D-galactose (12).³⁶ To access acceptor 13, glycosylation of a per-*O*-acetyl-β-GlcNAc with benzyl alcohol and ytterbium triflate,^{38,39} followed by deprotection under Zemplén conditions, gave the corresponding benzyl glycoside. A *tert*-butyldimethylsilyl protecting group was then introduced selectively on the primary hydroxyl to give the suitably protected glycosyl acceptor 13.⁴⁰ Regioselective glycosylation⁴¹ of 13 catalysed by BF₃ etherate gave β-1,4-linked disaccharides 14–18 containing the desired 6-modified galactose in 27–53% yield (Scheme 1).

Deprotection of compounds 14–17 by removal of the 6-*O*-silyl group, followed by de-*O*-acetylation under Zemplén conditions and removal of the anomeric benzyl protecting group by hydrogenolysis afforded target compounds 1–4. Azidodisaccharide 18 served as a vital divergent intermediate for amides 5–7. Staudinger reduction⁴² followed by per-*O*-acetylation and deprotection steps led to acetamide 5 from compound 18, whilst compounds 6 and 7 were accessed by Staudinger reduction and formation of amides under Schotten–Baumann conditions.⁴³ In order to facilitate immobilisation of this disaccharide library on a neutravidin-coated SPR imaging chip, compounds 1–7 were chemically derivatised at their reducing ends with 6-biotinamido hexanoic acid hydrazide, giving biotinylated glycosyl hydrazides^{32,33} (1Bt–7Bt, Scheme 1) that were suitable for printing.

Table 1 Library of biotinylated glycans used in this study


Group name	Glycotope structure	Common name
B301	Galβ1-4GlcNAcβ1-3Galβ1-3GlcNAc	L-NLe ^c
B266	Galβ1-4[Fucα1-3]GlcNAc	Le ^x
B186	Galβ1-4(6OSO ₃)Glc	6SuLac
B115	Galβ1-4GlcNAcβ1-3Galβ1-4Glc	LNnT
B111	(Galβ1-4GlcNAcβ) ₂	Di-LN
B81	Galβ1-4GlcNAc	LN
B80	Galβ1-4Glc	Lac
B82	Galβ1-3GlcNAc	Le ^c
B156	Galβ1-3[Fucα1-4]GlcNAc	Le ^a
B277	Galβ1-3GlcNAcβ1-3Galβ1-4GlcNAc	Le ^c -LN
B180	GalNAcβ1-4GlcNAc	LDN
B299	GlcNAcβ1-3Galβ1-3GlcNAc	GN-Le ^c
B181	GlcNAcβ1-3Galβ1-4GlcNAc	GN-LN
B121	GlcNAc	GN
B114	GlcNAcβ1-3Galβ1-4Glc	LNT ₂
B197	[3OSO ₃]Galβ1-4Glc	3'SuLac
B187	[3OSO ₃]Galβ1-4(6OSO ₃)Glc	6,3'-di-SuLac
B185	(6OSO ₃)Galβ1-4Glc	6'SuLac
B298	Neu5Acα2-3[Neu5Acα2-3Galβ1-3GalNAcβ1-4]Galβ1-4Glc	GD ₄
B273	Neu5Acα2-3GalNAcβ1-4GlcNAc	3'SLDN
B204	Neu5Acα2-3[Galβ1-3GalNAcβ1-4]Galβ1-4GlcNAc	GM ₂ (NAc)
B202	Neu5Acα2-3[Galβ1-3GlcNAcβ1-4]Galβ1-4Glc	GM ₁
B194	Neu5Acα2-3[Galβ1-4GlcNAcβ1-3] ₃	3'SLN-LN-LN
B178	Neu5Acα2-3[Galβ1-4GlcNAcβ] ₂	3'-SLN-LN
B177	Neu5Acα2-3[GalNAcβ1-4]Galβ1-4Glc	GM ₂
B174	Neu5Acα2-3Galβ1-3[Fucα1-4]GlcNAc	SLe ^a
B157	Neu5Acα2-3Galβ1-4[Fucα1-3]GlcNAc	SLe ^x
B85	Neu5Acα2-3Galβ1-3GlcNAc	3'SLe ^c
B84	Neu5Acα2-3Galβ1-4GlcNAc	3'SLN
B83	Neu5Acα2-3Galβ1-4Glc	3'SLac
B93	Neu5Gcα2-6Galβ1-4GlcNAc	6'S(Gc)LN
B90	Neu5Gcα2-3Galβ1-4GlcNAc	3'S(Gc)LN
B89	Neu5Gcα2-3Galβ1-4Glc	3'S(Gc)Lac
B274	Neu5Acα2-6GalNAcβ1-4GlcNAc	6'SLDN
B179	Neu5Acα2-6[Galβ1-4GlcNAcβ] ₂	6'-SLN-LN
B87	Neu5Acα2-6Galβ1-4GlcNAc	6'SLN
B86	Neu5Acα2-6Galβ1-4Glc	6'SLac
B184	Neu5Acα2-8Neu5Acα2-3[GalNAcβ1-4]Galβ1-4Glc	GD ₂
B108	Neu5Acα2-8Neu5Acα2-8Neu5Acα2-3Galβ1-4Glc	GT ₃
B107	Neu5Acα2-8Neu5Acα2-3Galβ1-4Glc	GD ₃

SPR imaging analysis of RCA₁₂₀ binding to a 6-substituted galactoside library

In order to assess RCA₁₂₀ recognition of unnatural galactoside derivatives, this modified library (**1Bt**–**7Bt**) was printed directly onto an SPR imaging chip, as described earlier. Equilibrium binding data are reported as a bar chart for glycans printed at 100 μg mL⁻¹ concentration (*ca.* 150 μM) in Fig. 3.††

As was immediately apparent from Fig. 3, replacement of the 6-hydroxyl group of galactose by fluorine (**2Bt**), or its removal altogether (**3Bt**), had limited impact on RCA₁₂₀ binding (compared to **1Bt**). In contrast, removal of the 5-hydroxymethyl group, to create **4Bt**, essentially abolished binding. A significant increase (3–4 fold) in SPR responses was generated by the presence of the designed amides [6'-NHAc-(**5Bt**), 6'-NHBz-(**6Bt**) and

6'-NH.CO.C₆H₁₁-Galβ1-4GlcNAc (**7Bt**)), when compared to LacNAc (**1Bt**). These results are consistent with those from our initial screen of forty natural glycans, where 6'-sialyl LacNAc terminated glycans (notably Neu5Ac/Gc-α2-6Galβ1-4GlcNAc types; red) showed significant increased affinities. However, these unnatural ligands represent a dramatic further increase and are the most potent known ligands thus far developed, to the best of our knowledge.

STD NMR spectroscopy analysis of RCA₁₂₀ binding to 6-modified galactosides

The data presented above clearly demonstrate that both terminal galactose and 6-modified galactose can act as motifs in ligands

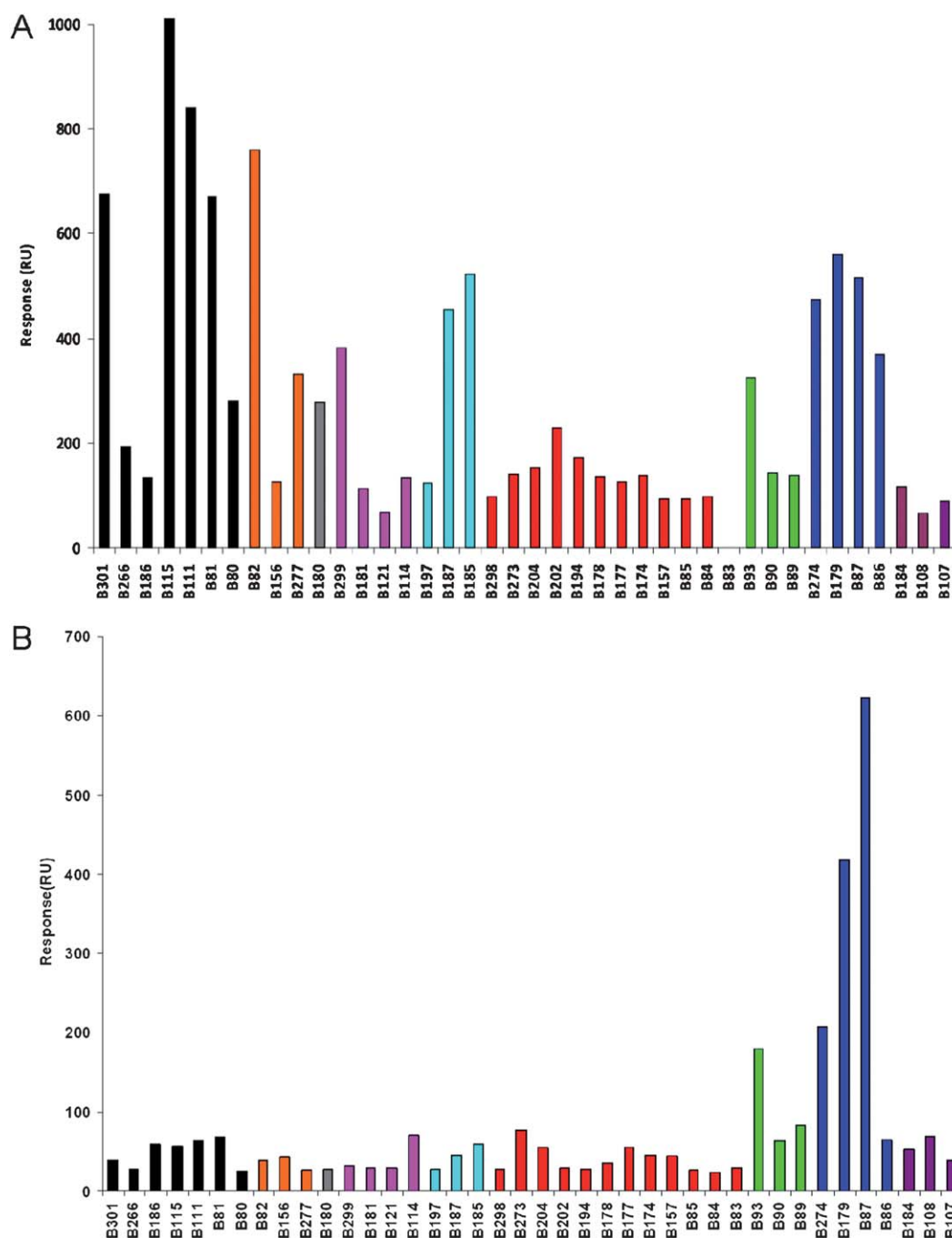


Fig. 1 SPR imaging responses for equilibrium binding of (A) RCA₁₂₀ (800 nM) and (B) SNA (80 nM) to an array of 40 biotinylated natural glycans (Table 1). Data errors were typically in the ± 10 –15% range.¹³

for RCA₁₂₀. What these SPRI glycoarray data are unable to confirm is whether both classes of ligand bind at the same site on the lectin. We addressed this critical point through a solution assay based on STD NMR spectroscopy.⁴⁴ STD NMR relies on saturation transfer from the lectin to the bound fraction of ligand upon lectin irradiation at a selected resonance, followed by spin diffusion.^{45,46} This approach is finding increasing use in the analysis of carbohydrate–protein interactions.^{47,48} Upon addition of Gal β 1-4Glc (1 mM) and the 6'-modified equivalent Neu5Ac α 2-6Gal β 1-4Glc (1 mM) as ligands in excess to individual solutions of RCA₁₂₀ (30 μ M), the STD NMR spectra were

recorded. Fig. 4A and 4B show the overlay of the 1D NMR reference spectra (in blue) and the corresponding STD NMR spectra (in red).

For Gal β 1-4Glc (Fig. 4A), the saturation was transferred mainly onto H2'-, H3'- and H4'-Gal (100, 78 and 72%) and to a lesser extent to the other protons (H1'-Gal 37% [not shown in Fig], H5'-Gal 41% and H6_{a,b}'-Gal 39 and 46%), consistent with literature data for methyl galactopyranoside.⁴³ This suggested that H2'-, H3'- and H4'-Gal were in close contact with the RCA₁₂₀ carbohydrate binding site. In the case of the 6'-modified equivalent Neu5Ac α 2-6Gal β 1-4Glc (Fig. 4B), H2'- and H3'-Gal

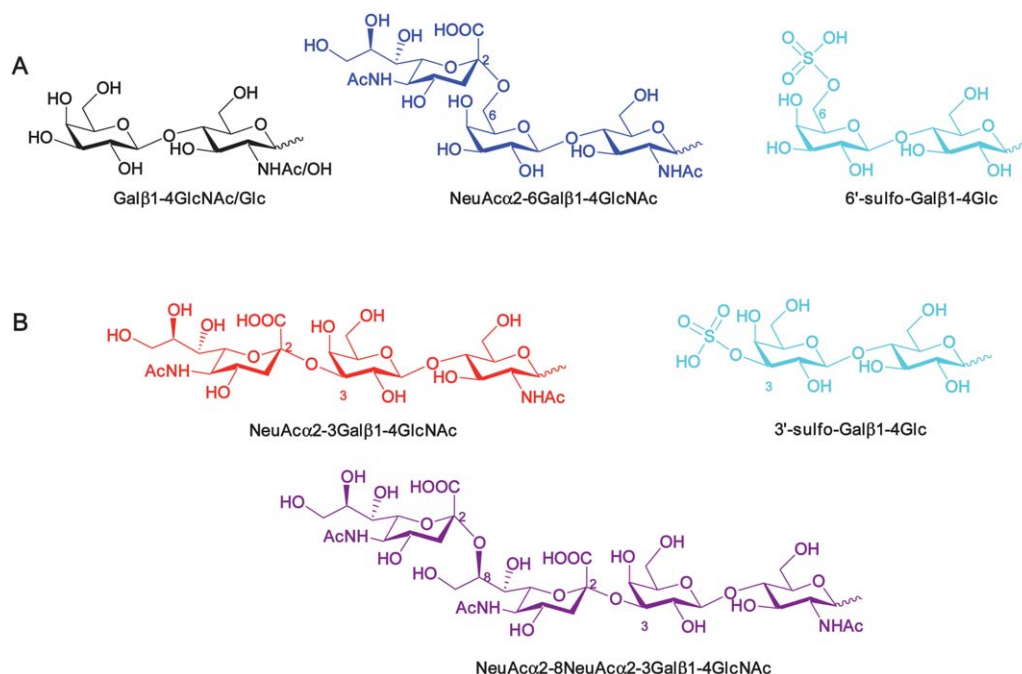
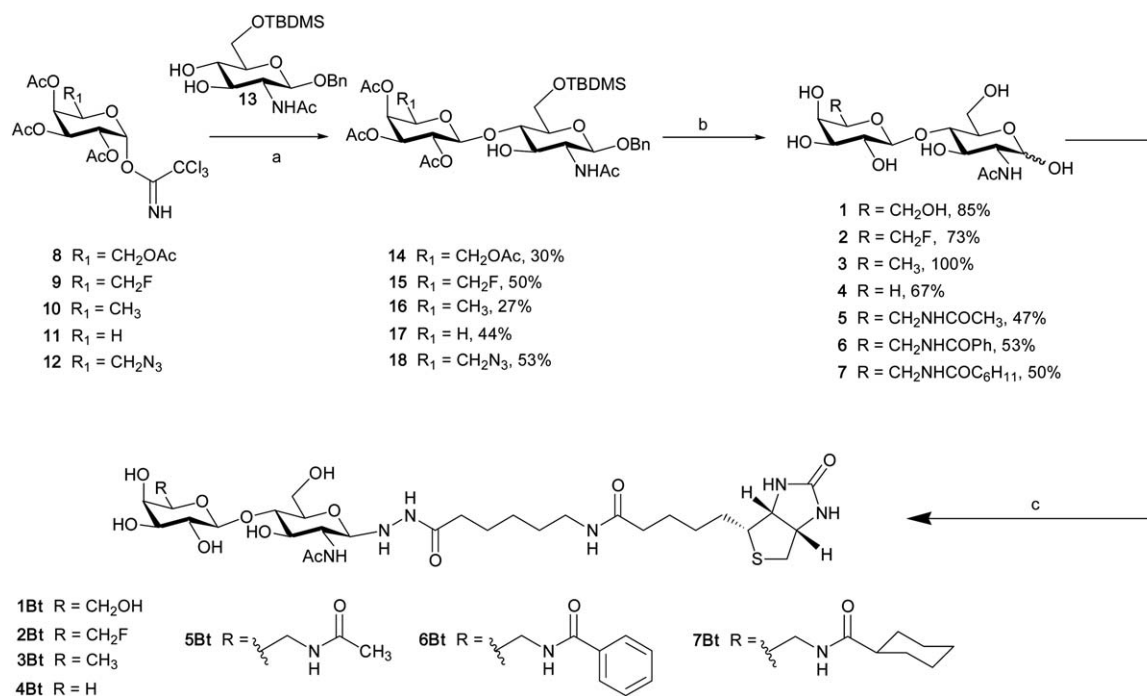


Fig. 2 Structures of representative oligosaccharides identified from SPRi array analysis that (A) serve as ligands for RCA₁₂₀ or (B) do not bind RCA₁₂₀. Colour coding refers to Table 1/ Fig. 1.

maintained the highest levels of saturation (100 and 72%, respectively), suggesting that these positions were integral to ligand recognition. Agreement with this observation can be found in lack of binding to RCA₁₂₀ of α2,3Neu5Ac-terminating oligosaccharides (red bars, Fig. 1) - *i.e.* substitution of the 3-position of galactose precludes binding to RCA₁₂₀, consistent

with the 3 position of galactose being situated in close proximity to the protein on binding to RCA₁₂₀. Significant differences were observed for saturation transfer for H4'-Gal signals in Galβ1-4Glc (72%) and Neu5Acα2-6Galβ1-4Glc (51%) and, to a lesser extent, for H5'-Gal (41% in Galβ1-4Glc; 51% in Neu5Acα2-6Galβ1-4Glc), reflecting changes in the network of interactions



Scheme 1 Reagents and conditions: a) $\text{BF}_3 \cdot \text{OEt}_2$, DCM, -40°C ; b) 1. $\text{BF}_3 \cdot \text{OEt}_2$, CH_3CN . 2. MeONa , MeOH. 3. H_2 , $\text{Pd}(\text{OH})_2/\text{C}$, MeOH for target compounds 1–4; b) 1. PPh_3 , H_2O ; 2. Ac_2O , Py; 3. $\text{BF}_3 \cdot \text{OEt}_2$; 4. MeONa , MeOH; 5. H_2 , $\text{Pd}(\text{OH})_2/\text{C}$, MeOH for target compound 5; b) 1. $\text{BF}_3 \cdot \text{OEt}_2$; 2. MeONa , MeOH; 3. PPh_3 , H_2O ; 4. RCOCl , aq. NaHCO_3 ; 5. H_2 , $\text{Pd}(\text{OH})_2/\text{C}$, MeOH for target compounds 6–7; c) 0.5 mol eq 6-biotinamido hexanoic acid hydrazide, MeOH/ H_2O / AcOH 92 : 6 : 2, 40°C , 24 h.

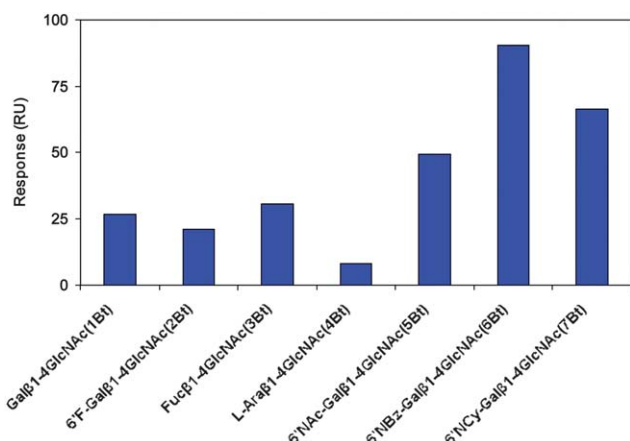


Fig. 3 SPR imaging responses for equilibrium binding of RCA₁₂₀ (800 nM) to an array of biotinylated Galβ1, 4-GlcNAc derivatives modified at the galactose 6-position (**1Bt–7Bt**). Data errors were typically in the ±10–15% range.¹³

which stabilises the ligand into the binding sites, particularly around the 4 and 5 positions of galactose. The protons from the added 6'-NeuAc group (H4'' to H9'') were also subject to some saturation transfer (38–55%, Fig. 4B), suggesting proximity to the protein surface in the RCA₁₂₀ complex.

Importantly, competition experiments using STD NMR with RCA₁₂₀ in the presence of both Galβ1-4Glc and Neu5Acα2-6Galβ1-4Glc in various ratios (SI, Figs. S18–S21†) provided strong support for these two ligands adopting very similar binding modes and hence competing for the same binding site on RCA₁₂₀.

Binding of 6-modified galactosides to RCA₁₂₀ in other interaction studies

Despite the widespread notion that RCA₁₂₀ is specific for non-reducing terminal β-linked D-galactose, numerous literature reports, employing a variety of techniques, give inconsistent evidence for the recognition of substituted galactosides. Early studies on glycan recognition reported that, α2-3Neu5Ac and α2-6Neu5Ac are “acceptable substituents on the terminal β-galactosyl moiety” for binding to ricin and its surrogate RCA₁₂₀.⁴⁹ However, while more recent observations by Gildersleeve and co-authors,⁵⁰ using a fluorescence glycoarray approach, identified Neu5Acα2-6Gal structures as RCA₁₂₀ binders (binding potency around 5 fold lower than lactose), α2-3Neu5Ac-containing glycans were not recognised in this assay. Data from frontal affinity chromatography studies using pyridylaminated (PA) oligosaccharides⁵¹ show that binding affinities of LacNAc-PA (K_D 4.1 μM) and 6'-sialyl LacNAc-PA (K_D 5.6 μM) to RCA₁₂₀ were comparable. Conversely, studies carried out using ELISA reported very low, if any, binding of sialylated lactosides to RCA₁₂₀.⁵² Fluorescence glycan array data from the Consortium for Functional Glycomics⁵³ using polyacrylamide-based glycoconjugates⁵⁴ show high RCA₁₂₀ binding to non-reducing terminal Gal, Galβ1-4Glc and Galβ1-4GlcNAc. Interestingly, some but not all oligosaccharides terminating in the Neu5Acα2-6Galβ1-4GlcNAc sequence were also strongly recognised on these arrays (see SI, Fig. S17†), but glycans terminating in α2,3-Neu5Ac or α2,8-diNeu5Ac were not recognised. The SPRi data reported in the present study are in agreement with the Gildersleeve data, demonstrating binding to α2,6-Neu5Ac-Gal but not to α2,3-Neu5Ac-Gal (blue and red in Fig. 1, respectively). Can these conclusions be rationalised on a structural basis?

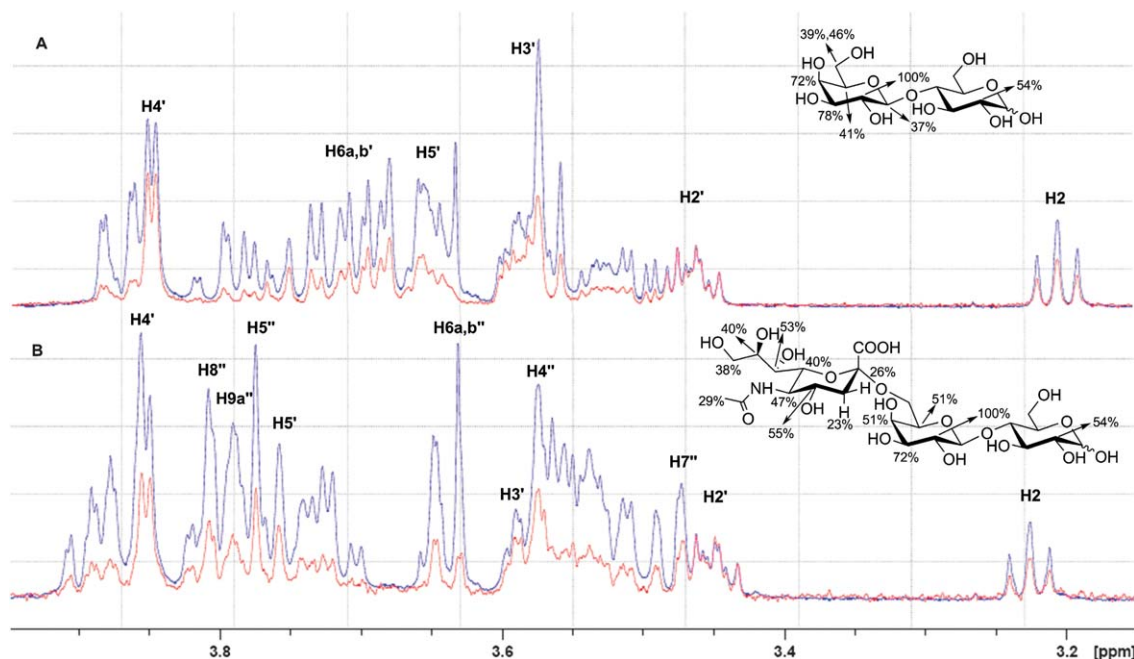


Fig. 4 The interaction of (A) Galβ1-4Glc (Lac, 1 mM) and (B) Neu5Acα2-6Galβ1-4Glc (6'SL, 1 mM) with RCA₁₂₀ (30 μM) by STD-NMR. The reference 1D NMR spectra (blue) are overlaid with the corresponding STD NMR spectra (red). The extent of saturation is expressed as relative intensity of the signals, normalised to the highest intensity signal (H2'-Gal for both (A) and (B)), to determine the binding epitopes. Gal residue protons are labelled as H_x', whilst Neu5Ac protons are labelled as H_x'. The chemical structures report the extent (%) of saturation for convenience.

Structural analysis of the binding of ricin and RCA₁₂₀ with oligosaccharide ligands

The X-ray crystal structure of RCA₁₂₀ bound to β -D-galactopyranose⁵⁵ reveals that each molecule of the agglutinin incorporates one carbohydrate binding site capable of forming four hydrogen bonds with the β -Gal ligand: between Lys40 and OH-2, Asn46 and OH-3, Gly25 and OH-1, Asp22 and OH-4 (Fig. 5a). A hydrogen bonding network is further stabilised by hydrophobic interactions between Trp37 and the hydrophobic lower face of β -Gal (Fig. 5b). Overall, the crystallographic data clearly illustrate that the galactose 6-OH group does not make key polar contacts with the protein and suggest that substitution at this position may be accommodated; these observations are critically consistent with the data that we have determined in the present study.

This conclusion is further supported by low-energy conformer distribution analyses and nOe NMR studies^{56,57} on ricin and RCA₁₂₀ using a series of deoxy-sugar derivatives of methyl β -lactoside. These studies have also highlighted the importance of positions 2, 3 and 4 in D-galactoside ligands as key polar groups for the interaction. Furthermore, STD NMR studies⁴⁴ show saturation transfer from RCA₁₂₀ to H-2, H-3 and H-4 (87, 100 and 87%, respectively) of methyl β -galactoside, but not to H-6, which is consistent with scope to accommodate modification of the galactose 6-OH group. Again, consideration of structural data supports the notion that alteration and substitution of the 6-position of galactose can be tolerated by RCA₁₂₀.

Conclusions

The need for fast and sensitive detection methods for the unequivocal identification of ricin led us to explore SPR imaging of glycoarrays for screening ligands of the ricin surrogate RCA₁₂₀. SPRI analysis of RCA₁₂₀ interaction with a 40 compound glycoarray highlighted the potential of 6-modified D-galactosides to serve as ligand for this lectin; these observations were further substantiated by STD NMR spectroscopy. Non-natural 6-modified galactosides inspired by these observations were designed, synthesised and subsequently shown to exhibit up to a 3–4 fold enhanced RCA₁₂₀ binding compared to the parent galactoside. To our knowledge, these are the most potent non-natural small molecule ligands currently known for RCA₁₂₀. In

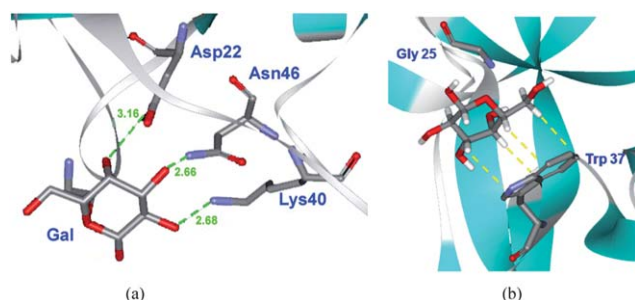


Fig. 5 (a) Interaction of galactose (Gal) hydroxyl groups with amino acid (AA) residues of RCA₁₂₀. In green, H-bond (Gal–OH \cdots NH₂–AA) distances expressed in Å (OH-2: 2.68, OH-3: 2.66, OH-4: 3.16); (b) In yellow, hydrophobic interactions between the lower face of Gal and Trp37 of RCA₁₂₀ (PDB code 1RZO).⁵⁵

summary, this study demonstrates the utility of glycoarrays and SPRI for both identifying and optimising carbohydrate ligands (Fig. 6) in the context of devising novel ligands for ricin sensor applications.

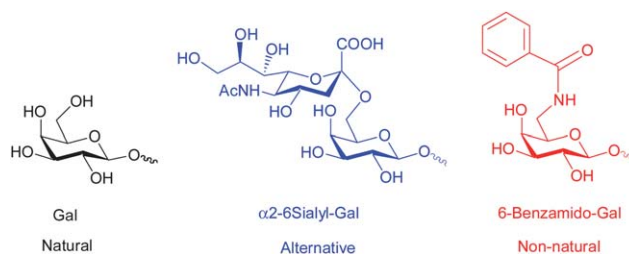


Fig. 6 RCA₁₂₀ ligands explored and developed in this study.

Experimental

Carbohydrate microarrays were prepared using biotinylated glycans, methods and hardware essentially as reported previously.¹² Along with details of the synthesis, characterisation of the modified *N*-acetylglucosamines, protocols for the biotinylation of these reducing sugars and STD NMR competition experiments, this information can be found in the Electronic Supporting Information. All STD NMR experiments were performed on a Bruker Avance II 600 MHz spectrometer equipped with a TCI cryoprobe; spectra were processed on Bruker Topspin 2.1 software. A Bruker pulsed sequence for STD-NMR (Bruker pulse sequence code: stddiffesp.3) was used with a shaped pulse train for saturation alternating between on and off resonance and a spoil pulse to destroy unwanted magnetization. The sequence includes solvent suppression using excitation sculpting with gradients and spinlock to suppress protein signals. On resonance was set at 360 Hz, whilst off resonance was set at $-50,000$ Hz. The number of scans varied with sample (typically 256). NMR samples were prepared in 500 μ L of 99.9% D₂O buffer containing 10 mM phosphate buffer, 150 mM NaCl at pH 7.4 (not corrected for D₂O). Solute exchange was achieved by ultrafiltration of the RCA₁₂₀ sample with Amicon® Ultra (Millipore) centrifugal filters (membrane cut-off 30 kDa). Addition of the saccharides Gal β 1-4Glc and Neu5Ac α 2-6Gal β 1-4Glc (Carbosynth Limited) to the protein NMR sample was from concentrated stock solutions in 99.9% buffered D₂O. The ratio of protein to oligosaccharide was typically *ca.* 1 : 33. In competition experiments, the competing carbohydrate ligand was in 4-fold excess with respect to the primary carbohydrate ligand and therefore in a 1 : 132 ratio with the protein. As far as carbohydrate epitope mapping analysis was concerned, the STD integrals of the individual protons of the two disaccharides (Gal β 1,4Glc and Neu5Ac α 2,6Gal β 1,4Glc) were referenced to the strongest STD signal in each spectrum (H2'-Gal in both the saccharides), which was assigned a value of 100%.

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Notes and references

¶ Current analytical procedures for the detection of ricin are summarised in ref. 15.

|| Abbreviations: Gal (galactose), GalNAc (*N*-acetylgalactosamine), Glc (glucose), GlcNAc (*N*-acetylglucosamine), NeuAc-2,3 (α 2,3-linked *N*-acetylneuraminic acid), NeuGc-2,3 (α 2,3-linked *N*-glycolylneuraminic acid), NeuAc-2,6 (α 2,6-linked *N*-acetylneuraminic acid), NeuGc-2,6 (α 2,6-linked *N*-glycolylneuraminic acid) and NeuAc-2,8 (α 2,8-linked *N*-acetylneuraminic acid). Colloquially, *N*-acetylneuraminic acid is often simply referred to as sialic acid.

** The relative ranking of binding events are used throughout this study. Consistent with problems experienced elsewhere for quantifying protein–carbohydrate interactions by SPR,³⁰ and cautionary notes on the subject, direct fitting of SPR data to simple 1 : 1 or 2 : 1 binding models is not always possible, or appropriate.³¹

†† SPR signals obtained for printed glycosyl hydrazones interacting with lectins were routinely much lower than for the corresponding glycosides (compare Fig. 5 and Fig. 2, respectively). However, where investigated, the relative ranking of binding was independent of the nature of the linkage.

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